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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/535,736	05/18/2005	Junichi Inagawa	B 520	9426
22840	7590	12/01/2008	EXAMINER	
GE HEALTHCARE BIO-SCIENCES CORP.			HOBBS, LISA JOE	
PATENT DEPARTMENT				
800 CENTENNIAL AVENUE			ART UNIT	PAPER NUMBER
PISCATAWAY, NJ 08855			1657	
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			12/01/2008	PAPER

**Please find below and/or attached an Office communication concerning this application or proceeding.**

The time period for reply, if any, is set in the attached communication.

<b>Office Action Summary</b>	<b>Application No.</b>	<b>Applicant(s)</b>	
	10/535,736	INAGAWA ET AL.	
	<b>Examiner</b>	<b>Art Unit</b>	
	Lisa J. Hobbs	1657	

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

#### Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

#### Status

1) Responsive to communication(s) filed on 09 September 2008.

2a) This action is **FINAL**.                            2b) This action is non-final.

3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

#### Disposition of Claims

4) Claim(s) 1-15 and 18-21 is/are pending in the application.

4a) Of the above claim(s) \_\_\_\_\_ is/are withdrawn from consideration.

5) Claim(s) \_\_\_\_\_ is/are allowed.

6) Claim(s) 1-15 and 18-21 is/are rejected.

7) Claim(s) \_\_\_\_\_ is/are objected to.

8) Claim(s) \_\_\_\_\_ are subject to restriction and/or election requirement.

#### Application Papers

9) The specification is objected to by the Examiner.

10) The drawing(s) filed on \_\_\_\_\_ is/are: a) accepted or b) objected to by the Examiner.

Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).

Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).

11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

#### Priority under 35 U.S.C. § 119

12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).

a) All    b) Some \* c) None of:

1. Certified copies of the priority documents have been received.
2. Certified copies of the priority documents have been received in Application No. \_\_\_\_\_.
3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

\* See the attached detailed Office action for a list of the certified copies not received.

#### Attachment(s)

1) Notice of References Cited (PTO-892)

2) Notice of Draftsperson's Patent Drawing Review (PTO-948)

3) Information Disclosure Statement(s) (PTO/SB/08)  
Paper No(s)/Mail Date \_\_\_\_\_.

4) Interview Summary (PTO-413)  
Paper No(s)/Mail Date. \_\_\_\_\_.

5) Notice of Informal Patent Application

6) Other: \_\_\_\_\_.

## **DETAILED ACTION**

### ***Continued Examination Under 37 CFR 1.114***

A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after final rejection. Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, the finality of the previous Office action has been withdrawn pursuant to 37 CFR 1.114. Applicant's submission filed on 09 September 2008 has been entered.

### ***Claim Status***

Claims 1-15, 18-21 are active in the case. Claims 16-17 have been cancelled by amendment. Claims 1-15 and 18-21 are under examination

### ***Claim Rejections - 35 USC § 103***

The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later

invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

The factual inquiries set forth in *Graham v. John Deere Co.*, 383 U.S. 1, 148 USPQ 459 (1966), that are applied for establishing a background for determining obviousness under 35 U.S.C. 103(a) are summarized as follows:

1. Determining the scope and contents of the prior art.
2. Ascertaining the differences between the prior art and the claims at issue.
3. Resolving the level of ordinary skill in the pertinent art.
4. Considering objective evidence present in the application indicating obviousness or nonobviousness.

Claims 1-15 and 18-21 are rejected under 35 U.S.C. 103(a) as being unpatentable over Wagner et al. (WO 2001/72458) in view of Bosman et al. (WO 1999/00670), Barner et al. (US 5986066), Badley et al. (US 6294391) and Nelson et al. (US 5955729). Wagner et al. teach heterofunctional cross linking reagents, protein labeling reagents, protein conjugates and their compositions, support-bound cross linking groups, modified supports and protein arrays for site specific binding of proteins, they teach techniques for attaching a biomolecule (a protein) containing a tag by binding sites for the biomolecule tag and for covalently attaching a biomolecule to activated reactive groups (support-bound cross linking groups) to a solid support. Barner et al. teach cross-linking molecules, which molecules will biologically recognize target molecules, to a solid phase using carrier molecules. Bosman et al. teach methods of covalently immobilizing biomolecules by means of a His-tag and using a substrate that biologically recognizes the His tag. Badley et al. teach methods of detecting the presence of an analyte of interest in a sample, the method comprising the steps of: providing a binding partner reversibly immobilized on a solid support, said binding partner having binding specificity for the analyte;

contacting the sample with the solid support; specifically displacing the binding partner from the solid support in response to the presence of the analyte of interest in the sample, said displacement causing a reduction in the mass of material immobilized on the solid support, thereby generating a detectable change in a mass-dependent property of the solid support; and detecting said change, while Nelson et al. teach detection of analytes using surface plasmon resonance.

Wagner et al. teach that it is known to attach a protein to a solid support by associating a protein containing a tag with a protein tag binder, see page 6, lines 3-10 (claims 1, 12, 18, and 20); they also disclose a method for covalently attaching a protein to the surface by linking groups (claim 2). Wagner et al. also teach the use of an amino group from the biomolecule and a carboxyl group of the sensor chip to create a covalent bond (claim 3). As well, Barner et al. explicitly state in columns 3 and 4 that reactive functional groups, such as COOH or NH<sub>2</sub> are well-known for use as covalent attachment points for immobilizing biomolecules (claims 2-3).

Additionally, Wagner et al. teach naturally binding molecules (claim 11), such as antigen/antibody recognition epitopes, as protein-tag binders. On page 13 they teach His tags (claims 9-10). However, they do not teach details of His tag antibody and antigen reactions. Bosman et al. teach detailed methods of using His tags and His tag antibodies, see entire document, including the complexing of glycoproteins to metal affinity resin on page 3 (claim 19).

At page 26, Wagner et al. teach the introduction of introduce histidine tags into the protein (claim 4) and then the binding of the protein to a sensor chip coated with nitrilotriacetic acid (NTA) through Ni<sup>2+</sup> (claims 4-7). Wagner et al. do not teach the use of iminodiacetic acid

(IDA), however it is taught by Bosman et al., page 3, that IDA/Ni<sup>2+</sup> can be used as an alternative to NTA in His tag immobilization (claim 8).

Wagner et al. do not specifically teach low molecular weight compound binding (claims 13 and 15), however they, Barner et al., and Bosman et al., do describe multiple binding substituents, including a statement by Bosman et al. that the invention is “to simultaneously provide a universal detection method for biomolecules that contain a His tag”, page 5, and Wagner et al. teach protein-protein, protein-nucleic acid, protein-drug, and protein-ligand interactions, see page 2, which encompasses a large range of molecular weights.

The use of surface plasmon resonance to measure and detect biomolecules and analytes of interest (claim 14) is known in the art, as described by Nelson et al., see entire document, while Badley et al. specifically teach methods of detecting the presence of an analyte of interest in a sample, the method comprising the steps of: providing a binding partner reversibly immobilized on a solid support, said binding partner having binding specificity for the analyte; contacting the sample with the solid support; specifically displacing the binding partner from the solid support in response to the presence of the analyte of interest in the sample, said displacement causing a reduction in the mass of material immobilized on the solid support, thereby generating a detectable change in a mass-dependent property of the solid support; and detecting said change, paragraph 23, using “a number of mass-dependent properties which can be detected, for example, by acoustic wave or evanescent wave type sensors, or by surface plasmon resonance (SPR) detectors, all of which are known in the art”.

It would have been obvious to one of ordinary skill in the art at the time the invention was made to combine the teachings of Wagner et al. with Barner et al., Bosman, et al., Badley et

al., and Nelson et al. in order to achieve the invention as claimed in the claims under examination. As demonstrated above, methods for the immobilization of biomolecules involving covalent binding of substituents by chemical groups or by antibody/antigen binding, assisted by other chemicals, using components naturally present or added to the molecules as needed, were known in the art and the claims, as presented, are rendered obvious.

***Response to Arguments***

Applicant's arguments filed 15 July 2008 have been fully considered but they are not persuasive. Applicant argues that the references cited neither teach nor suggest a method of immobilizing a biomolecule to a substrate having both tag binding sites and activated reactive groups. However, Bosman et al. discuss the use of His tags in immobilization of biomolecules and the specifics of using tags with and without additional covalent binding to the support. They discuss a method "wherein the presence of His-tags is exploited for covalent immobilisation of a biomolecule that contains said His-tag, and wherein the amino acid residues that comprise said His-tag are directly involved in the covalent bond" (page 5). As well, they describe methods to increase the probability of a reaction with the his residues of the His-tag of the biomolecule, while increasing the reaction of the membrane or carrier to which said biomolecule has to be immomobilised covalently (page 9). Finally, on page 12, they discuss that biomolecules that use His-tags as crosslinking elements as well as the biomolecule being involved in a covalent immobilization forming an aggregate. On page 13, they specifically describe the inclusion of methods for covalent linkage between two biomolecules, of which at least one, or sometimes both, contains a His-tag.

Applicants particularly argue that Bosman et al. do not specifically teach that the second form of binding occurs specifically between non-tag elements of the construct. This argument has been considered but is not persuasive. Claim 1 recites that the biomolecules of interest have at least one tag which will bind to the immobilization substrate and also have "reactive groups which are **capable of** forming covalent bonds with the non-tag part of the biomolecule" (emphasis added). As well, the subsequent dependent claims, particularly claim 2, do not recite positively that the second form of binding between the molecule of interest and the substrate is not related to the tag. Claim 1 recites that there are reactive groups that are capable of forming a bond without the tag. Claim 2 recites that the first step is that reactive groups are activated, but does not recite that these are not reactive groups which do not bind to the tag (claim 1 merely recites that the reactive groups are "capable of" forming bonds with parts of the molecule other than the tag). Claim 2 then recites that the biomolecule is immobilized on the substrate through a tag interaction and a covalent bond, but again does not state a positive requirement that the covalent bond is not through the tag. As well, none of claims 9, 13, or 18 recite positively that the second bond is not related to the tag at all, which would clarify the statement "capable of" in the independent claim, claim 1.

### ***Conclusion***

No claims are allowed.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Lisa J. Hobbs whose telephone number is 571-272-3373. The examiner can normally be reached on Monday to Friday, 8:00 a.m. to 4:30 p.m.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Jon P. Weber can be reached on 571-272-0925. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

/Lisa J. Hobbs/  
Primary Examiner  
Art Unit 1657

ljh